



Standard Guide for Preparation of Aluminum Alloys for Electroplating¹

This standard is issued under the fixed designation B 253; the number immediately following the designation indicates the year of original adoption or, in the case of revision, the year of last revision. A number in parentheses indicates the year of last reapproval. A superscript epsilon (ε) indicates an editorial change since the last revision or reapproval.

This standard has been approved for use by agencies of the Department of Defense.

ε¹ NOTE—The Warning notes in 7.1 and X1.8 were updated in October 2005.

1. Scope

1.1 This guide covers cleaning and conditioning treatments used before metal deposition (Section 5), and immersion deposit/strike procedures (Section 6) that enhance the adhesion of metals that are subsequently applied to aluminum products by electrodeposition or by autocatalytic chemical reduction.

1.2 The following immersion deposit/strike procedures are covered:

1.2.1 Zinc immersion with optional copper strike (6.3).

1.2.2 Zinc immersion with neutral nickel strike (6.4).

1.2.3 Zinc immersion with acetate-buffered, nickel glycolate strike (6.5).

1.2.4 Zinc immersion with acid or alkaline electroless nickel strike.

1.2.5 Tin immersion with bronze strike (6.6).

1.3 From the processing point of view, these procedures are expected to give deposits on aluminum alloys that are approximately equivalent with respect to adherence. Corrosion performance is affected by many factors, however, including the procedure used to prepare the aluminum alloy for electroplating.

1.4 This guide is intended to aid electroplaters in preparing aluminum and its alloys for electroplating.

1.5 *This standard does not purport to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this standard to establish appropriate safety and health practices and determine the applicability of regulatory limitations prior to use.* For specific precautionary statements see Section 7 and Appendix.

2. Referenced Documents

2.1 *ASTM Standards:*²

B 85 Specification for Aluminum-Alloy Die Castings

B 179 Specification for Aluminum Alloys in Ingot and Molten Forms for Castings from All Casting Processes

B 209 Specification for Aluminum and Aluminum-Alloy Sheet and Plate

B 209M Specification for Aluminum and Aluminum-Alloy Sheet and Plate [Metric]

B 221 Specification for Aluminum and Aluminum-Alloy Extruded Bars, Rods, Wire, Profiles, and Tubes

B 221M Specification for Aluminum and Aluminum-Alloy Extruded Bars, Rods, Wire, Profiles, and Tubes [Metric]

B 322 Guide for Cleaning Metals Prior to Electroplating

B 431 Practice for Processing of Mandrels for Electroforming³

E 527 Practice for Numbering Metals and Alloys (UNS)

3. Significance and Use

3.1 Various metals are deposited on aluminum alloys to obtain a decorative or engineering finish. The electroplates applied are usually chromium, nickel, copper, brass, silver, tin, lead, cadmium, zinc, gold, and combinations of these. Silver, tin, or gold is applied to electrical equipment to decrease contact resistance or to improve surface conductivity; brass, copper, nickel, or tin for assembly by soft soldering; chromium to reduce friction and obtain increased resistance to wear; zinc for threaded parts where organic lubricants are not permissible; tin or lead is frequently employed to reduce friction on bearing

¹ This guide is under the jurisdiction of ASTM Committee B08 on Metallic and Inorganic Coatings and is the direct responsibility of Subcommittee B08.02 on Pre Treatment.

Current edition approved Sept. 15, 2005. Published October 2005. Originally approved in 1951. Last previous edition approved in 1999 as B 253 – 87 (1999)^{ε1}.

² For referenced ASTM standards, visit the ASTM website, www.astm.org, or contact ASTM Customer Service at service@astm.org. For *Annual Book of ASTM Standards* volume information, refer to the standard's Document Summary page on the ASTM website.

³ Withdrawn.

surfaces. Nickel plus chromium or copper plus nickel plus chromium is used in decorative applications. Nickel plus brass plus lacquer or copper plus nickel plus brass plus lacquer is also used for decorative finishes, sometimes with the brass oxidized and relieved in various ways.

3.1.1 Electroless nickel may be applied as a barrier layer prior to other deposits, or for engineering purposes.

3.2 The preparation of aluminum and aluminum alloy mandrels for electroforming is described in Practice B 431.

4. Nature of Aluminum and Its Influence on Preparation

4.1 *Microstructure*—It is difficult to find a preplating procedure that is equally satisfactory for all types and tempers of aluminum alloys because the various alloys and products behave differently electrochemically due to their different compositions and metallurgical structures. When elements are added for alloying purposes, they may appear in an aluminum alloy in several different forms: that is, they may be in solid solution in the aluminum lattice, be present as microparticles of the elements themselves, or be present as particles of intermetallic compounds formed by combination with the aluminum. The several solid solution matrices and the 20 or more microconstituents that may occur in commercial alloys may have different chemical reactivities and electropotentials and their surfaces may not respond uniformly to various chemical and electrochemical treatments. In addition, the response may be influenced by variations in the microstructure of different lots of products of the same alloy. In some cases, these variations may be introduced or aggravated by preparation processes; for example, the heat generated in buffing. The electroplater needs to know the aluminum alloy that is to be processed in order to select the best electroplating procedure. In the absence of this information, there are so-called universal procedures that may be used. However these will not necessarily be the best or the most economical procedures for the alloy.

4.2 *Oxide Film*—In addition to differences in microstructure that may affect response to preplating treatments, all aluminum products have an ever-present natural oxide film. This oxide film can be removed by various acid and alkaline treatments and even though it reforms immediately on contact with aqueous solutions or air, it then is usually thinner and more uniform than the original film. The newly formed oxide film provides a more suitable surface for deposition of the first metallic layer.

5. Cleaning and Conditioning Treatments

5.1 To obtain consistent results for electroplating on aluminum alloys, it is essential that the various cleaning and conditioning treatments provide a surface of uniform activity for the deposition of the initial metallic layer. First, the surface should be free of any oil, grease, buffing compound, or other foreign material. For removing oil, grease, or buffing compound, use vapor degreasing,⁴ solvent washing, or solvent

emulsion cleaning. For removing buffing compound, specially formulated detergent type or modified detergent type buffing compound removers may also be used. If the deposits of soil are relatively light and fairly uniformly distributed, a mild etching type cleaner may also be used. A convenient one is a hot, aqueous carbonate-phosphate solution (X1.1). Other types of cleaners are used; for example, mildly alkaline or acidic soak cleaners are used to remove gross soils. Also available are a wide range of proprietary cleaners of the “non-etching” type. Some of these are actually buffered mixtures, similar to the carbonate-phosphate mixture (X1.1) where the so-called non-etching characteristics are obtained by buffering the solution to pH levels where the etching action becomes minimal. Others are truly non-etching types where etching is prevented by using silicate inhibitors, such as sodium metasilicate (Na₂SiO₃). These inhibitors always leave a film of aluminum silicate on the surface. When these materials are used, subsequent deoxidizing solutions should contain controlled amounts of fluoride salts to insure complete removal of the film.

NOTE 1—General information on the cleaning of metals is given in Guide B 322.

5.2 After cleaning, a conditioning treatment of the surface is generally required. For this to be effective, it must accomplish two things: (1) remove the original oxide film and (2) remove any microconstituents that may interfere with the formation of a continuous deposited metallic layer or that may react with subsequent electroplating solutions.

5.2.1 An effective conditioning treatment is immersion of the work in a warm sodium hydroxide solution (Appendix X1.3) followed by water rinsing and immersion in a nitric acid-bifluoride desmutting solution (Appendix X1.4). An alternative desmutting solution is sulfuric acid-hydrogen peroxide (Appendix X1.5).

NOTE 2—When an unmodified sodium hydroxide solution is used, etching may become nonuniform and heavy concrete-like scales may form on tank walls and heating surfaces, their development becoming more rapid as the concentration of dissolved aluminum increases. The incorporation of controlled amounts of deflocculating complexors such as sodium gluconate, sodium glucoheptonate, certain sugar derivatives, and certain substituted sugar amines will eliminate this problem. Many proprietary etching materials are so modified.

NOTE 3—The universal acid mixture (Appendix X1.9) is applicable to almost all alloys, and is especially desirable for use with alloys containing magnesium.

5.2.2 For heat-treated alloys (alloys in a “T” temper), it is important to remove the relatively thick, heat-treated oxide film before proceeding with subsequent conditioning treatments. Normally, heat-treated films are removed by machining, or by the polishing action on metal surfaces that are buffed.

5.2.2.1 In the absence of machining or buffing, controlled abrasive blasting may be used to remove this oxide. Fine abrasives such as aluminum oxide, ceramic beads, or glass beads may be used. Silicon carbide abrasives should be avoided. If aluminum oxide, or glass beads are used, subsequent treatments should include the use of an acid fluoride to ensure that any embedded aluminum oxide or silica is removed. However, surfaces of heat-treated alloys that are not machined or buffed should have the heat-treated film removed

⁴ For details on the proper operation and safety precautions to be followed in vapor degreasing, see *Handbook of Vapor Degreasing*, ASTM STP 310, ASTM, 1976.

with a deoxidizing etch to obtain uniform electroplating results. An effective deoxidizing etch is a hot sulfuric-chromic acid solution (Appendix X1.2). Suitable proprietary deoxidizing etches including some with no chromates are available. They should be used as recommended by the manufacturer.

5.2.3 For wrought alloys of the UNS A91100 and UNS A93003 types (see Specifications B 209 and B 209M) fairly good conditioning may be obtained by using the carbonate-phosphate cleaner (Appendix X1.1) followed by a nitric acid dip at room temperature (Appendix X1.6). These alloys do not contain interfering constituents and for some applications, this method of conditioning may be ample. If a silicate inhibited cleaner is used (see 5.1) the fluoride containing smut remover (Appendix X1.4) is preferred.

NOTE 4—In accordance with current ASTM practice and for international usage, the aluminum alloys have been classified in accordance with the Unified Numbering System (UNS) as detailed in Practice E 527 and listed in D556C.⁵

5.2.4 Another effective conditioning treatment for removing the surface oxide film and any undesirable microconstituents comprises the use of a hot sulfuric acid etch (Appendix X1.7). The time of the dip depends on the alloy involved. Generally the shorter time is used on castings. This treatment is satisfactory for all aluminum-magnesium alloys, both wrought and cast. It not only leaves the surface in an excellent condition for the deposition of the first metallic layer, but it also eliminates the undesirable effects of the magnesium-containing constituents in alloys of the UNS A95052, UNS A96061, and UNS A96063 types (see Specifications B 221 and B 221M).

5.3 The following are types of casting alloys containing high percentages of silicon: UNS A04130, UNS A14130, UNS A03800, (see Specification B 85), UNS A03561, and UNS A13560, (see Specification B 179). A dip at room temperature in a mixed acid solution (Appendix X1.8) containing nitric and hydrofluoric acids is recommended for conditioning the surface of these alloys. This treatment also removes the heat-treated film from unpolished, heat-treated castings.

6. Immersion Deposit/Strike Procedures

6.1 Following the cleaning and conditioning treatments, it is necessary to further treat the surface to obtain adequate adhesion of an electrodeposited metal on aluminum alloys. This section describes five commercially used procedures:

- 6.1.1 Zinc immersion with optional copper strike (6.3).
- 6.1.2 Zinc immersion with neutral nickel strike (6.4).
- 6.1.3 Zinc immersion with acetate buffered, nickel glycolate strike (6.5).
- 6.1.4 Zinc immersion with an acid or alkaline electroless nickel strike (6.6).
- 6.1.5 Tin immersion with bronze strike (6.7).

6.2 The immersion deposit/strike conditions recommended for each procedure give good results with many alloys of aluminum. However, some alloys and tempers may require slight modification of the processing conditions for best results.

6.3 Zinc Immersion with Optional Copper Strike:

6.3.1 In the zinc immersion step, the oxide film is removed from the surface to be electroplated and is replaced by a thin and adherent layer of metallic zinc. This provides a surface that responds to most of the electroplating procedures for plating other metals on zinc.

6.3.2 For the immersion step, a highly alkaline solution⁶ containing the following components can be used at room temperature (15 to 27°C).

Zinc Immersion Solution, Bath I	
Sodium hydroxide (commercial)	525 g/L
Zinc oxide (technical grade)	100 g/L

6.3.2.1 For best results, the sodium hydroxide must be low in sodium carbonate content (preferably under 2 % by weight) and the zinc oxide must be free of contamination.

NOTE 5—In the zinc immersion solutions in this standard, the purity of the ingredients often plays an important role in the successful operation of the process. This is particularly true of the zinc oxide used. Contamination of the zinc oxide with lead or arsenic can be especially troublesome. Proprietary, prepared powdered or liquid zincates are frequently used therefore, since they will have had all raw materials properly checked for purity.

6.3.2.2 The thickness and quality of the immersion film are influenced by the conditions of deposition. When deposition is too rapid, heavy, coarse, crystalline, and porous, non-adherent deposits are formed. Since the thinner zinc deposits give the best results, it is recommended that the temperature of the zincate solution be kept below 27°C and the immersion time be from 30 s to 1 min.

6.3.3 A modification of the basic zincate solution in most applications gives more uniform and satisfactory results. The modified zinc immersion procedure has the following advantages: (1) more uniform coverage by subsequent electroplating baths, (2) greater operating range for the “double immersion” version of the treatment (see 6.3.5), and (3) improved resistance to corrosion on all electroplated aluminum alloys except for the UNS A92024 and UNS A97075 alloys. The modified solution is prepared by dissolving the zinc oxide in a sodium hydroxide solution and cooling to room temperature. Before the bath is diluted to volume, a water solution of ferric chloride crystals and Rochelle salt (potassium sodium tartrate) is added. The bath should be stirred while the ferric chloride-Rochelle salt solution is added.⁷ The modified zincate solution is made up as follows:

Zinc Immersion Solution, Bath II	
Sodium hydroxide	525 g/L
Zinc oxide	100 g/L
Ferric chloride hexahydrate	1.0 g/L
Rochelle salt	10 g/L

6.3.3.1 This bath should also be operated under 27°C and for immersion times of the order of 30 s to 1 min. It is recommended that Bath II be utilized whenever the “double immersion” treatment is employed. Likewise, it will be found

⁵ D556C Metals and Alloys in the Unified Numbering System, available from ASTM Headquarters. Order PCN 05-0564-02.

⁶ Sodium zincate solutions of this general type are now being replaced by newer modified zincate compositions.

⁷ There are proprietary zincate solutions available containing cations other than iron (also various other additions such as complexing agents or chelating agents or both). A solution containing copper and nickel, as well as zinc, is described by Schaer, G., *Plating and Surface Finishing*, 68,51 (March 1981).

advantageous on all wrought and cast alloys, except the UNS A92024 and UNS A97075 types, for corrosion-resistant applications.

6.3.3.2 With both of the solutions (Baths I and II), the rinse immediately after the zinc immersion step is critical. The activity of the solution increases rapidly with dilution. Because of the high concentrations used, the solution is viscous. If this viscous layer is not promptly removed in the rinsing step, the diluted film may deposit a loose, spongy zinc film in the rinse, thereby destroying an otherwise acceptable zinc film. Therefore, rinses must be strongly agitated so that this film is rapidly and uniformly removed. Spray rinsing at moderate to high pressure is preferred where the part configuration is such that the sprays can impinge on all surfaces.

6.3.4 Dilute versions of the modified zinc immersion procedures⁷ have been developed for applications where rinsing and drag-out are problems. The bath viscosity is reduced by lowering the concentration of the principal components. In using the dilute baths, a low film weight must be maintained by a closer control of operating conditions and by addition agents. Two typical dilute baths are prepared as follows:

Zinc Immersion Solution, Bath III	
Sodium hydroxide	50 g/L
Zinc oxide	5 g/L
Rochelle salt	50 g/L
Ferric chloride hexahydrate	2 g/L
Sodium nitrate	1 g/L
Zinc Immersion Solution, Bath IV	
Sodium hydroxide	120 g/L
Zinc oxide	20 g/L
Rochelle salt	50 g/L
Ferric chloride hexahydrate	2 g/L
Sodium nitrate	1 g/L

6.3.4.1 Bath IV will provide a much greater zinc reserve for high-production work with only a small sacrifice in rinsing and drag-out properties. When using these dilute solutions, the temperature must be maintained between 20 to 25°C and the immersion time must not exceed 30 s.

6.3.4.2 A more highly modified zincate (modified with copper, nickel, and iron) has been described by Wyszynski.⁸ It has much greater tolerance for variations in operating conditions, especially temperature and time of immersion, and permits processing a wider variety of alloys without resorting to the double zincating treatment (6.3.5). Because the quaternary alloy deposited by immersion is much less active than the relatively pure zinc from many immersion baths, subsequent electroplated deposits are applied with less difficulty.

6.3.5 A variation of the zincate treatment that has considerable merit consists of a double zinc immersion treatment with the first zinc layer being removed by a dip in a room temperature solution of 500 mL of concentrated, nitric acid (67 mass %, density 1.40 g/mL) diluted to 1 L. With this procedure, the first immersion dip removes the original oxide film and replaces it with a zinc layer. Removal of the zinc layer by the nitric acid dip leaves the surface in suitable condition for deposition of the final zinc immersion layer.

6.3.6 The concentrated zincate solutions (Baths I and II) are very viscous and losses occur largely from drag-out. This is advantageous as it limits the accumulation of impurities resulting from attack on the aluminum. It has the disadvantage however in that it increases the load on the waste disposal system.

6.3.7 The specific gravity of the concentrated solutions should be checked occasionally and any loss made up by adding more of the components. Loss of volume by dragout should be corrected by the addition of more solution of the specific composition. The dilute solutions (Baths III and IV and those recommended by Schaer and Wyszynski) should be controlled by chemical analysis of the caustic, zinc, and modifying metal concentration.

6.3.8 When a properly conditioned aluminum alloy article is immersed in the zincate solution, the thin natural oxide film that is present on the surface of the article dissolves and, as soon as underlying aluminum is exposed, it also starts to dissolve and is immediately replaced by an equivalent weight of zinc. When the aluminum surface is completely covered with an extremely thin layer of zinc, action in this solution virtually ceases.

6.3.9 With correct procedure, the resulting zinc deposit will be fairly uniform and firmly adherent to the surface. The appearance of the surface, however, will vary with the alloy being coated as well as the rate at which the coating forms. The weight of zinc deposit should be of the order 15 to 50 µg/cm², corresponding to a thickness of 20 to 70 nm. Generally, it is desirable to limit the weight of the deposit to not over 30 µg/cm². The thinner and more uniform zinc deposits are the most suitable for electroplating preparation and for the performance of electroplated coatings in service. Heavy zinc deposits tend to be spongy and less adherent and do not provide as good a surface for obtaining adherence as the thinner deposits. The weight of the zinc deposit will vary with the alloy and the conditioning treatment that is used.

6.3.10 After the surface of an aluminum alloy article has been conditioned and the zinc immersion deposit has been formed, other metals can be electroplated on this surface by any of the methods suitable for electroplating on zinc. Ordinarily, it is advisable to apply a suitable copper strike over the zinc-immersion layer before other metals are deposited. Silver, brass, zinc, nickel, or chromium, however, may be deposited on the zinc immersion layer provided the electroplating procedures are suitable for electroplating over zinc.

6.3.11 When a copper strike is to be used over the zinc immersion layer, a tartrate-type copper cyanide solution operated as follows is recommended:

Tartrate-Type Copper Strike Solution	
Copper cyanide	42.0 g/L
Total sodium cyanide	50.0 to 55.0 g/L
Sodium carbonate	30.0 g/L
Rochelle salt	60.0 g/L
Free sodium cyanide	5.5 to 10.5 g/L

The work is introduced with the electrical circuit connection made for "live" entry (cathodic).

Temperature	40 to 55°C
pH	10.2 to 10.5
Current density	260 A/m ²
Time	2 min

⁸ Wyszynski, A. E., et al, Transactions of the Institute of Metal Finishing, (England), Vol 45, 1967, pp. 147–154; Vol 59, 1981, pp. 17–24. Wyszynski, A. E., and Such, T. E., *Plating*, Vol 10, 1965, pp. 1027–1034.

6.3.11.1 Reduce cathode current density to 130 A/m² and electroplate for an addition 3 to 5 min.

6.3.12 After this strike, the work can be transferred to other standard electroplating solutions for further electroplating.

6.4 Zinc Immersion/Neutral Nickel Strike:⁹

6.4.1 Aluminum parts with cleaned and conditioned surfaces are given a double zinc immersion treatment as described in 6.3.3 and 6.3.5. Recommended times for the first and second zinc immersion are 45 s and 30 s, respectively.^{10,11,12}

6.4.2 After water rinsing, the zincated parts are given a nickel strike as follows:

6.4.2.1 The power source should be on and the electrical circuit connection made for “live” entry before immersing the work in the strike electrolyte.

6.4.2.2 (GMR) Neutral Nickel Strike Treatment Electrolyte:

Nickel sulfate 7H ₂ O	142 g/L
Ammonium sulfate	34 g/L
Nickel chloride 6H ₂ O	30 g/L
Sodium citrate	140 g/L
Sodium gluconate	30 g/L
Temperature	57 to 66°C
pH at 60°C	6.8 to 7.2
Current density	950 to 1300 A/m ²
Time	30 to 45 s
Agitation	Non-air type

6.4.2.3 Reduce cathode current density to 400 to 550 A/m² and electroplate for an additional 3 to 5 min.

6.4.3 After receiving the above neutral nickel strike, the aluminum parts can be electroplated with other metals using standard electroplating solutions.

6.5 Zinc Immersion/Nickel Glycolate Strike:¹⁰

6.5.1 Aluminum parts with cleaned and conditioned surfaces are given a zinc immersion treatment as described in 6.3. A single or a double immersion treatment may be used.

6.5.2 After water rinsing, the zincated parts are given a nickel strike in the mildly acid electrolyte with the following process conditions:

	Acetate Buffered Nickel Glycolate Strike Treatment Electrolyte
Nickel acetate 4H ₂ O	65 g/L
Boric acid	45 g/L
70 % Glycolic acid	60 mL/L
Saccharin	1.5 g/L
Sodium acetate	50 g/L
Temperature	room
pH	5.5 to 6.8

⁹ US Patent 3,417,005 assigned to General Motors Corporation.

¹⁰ Missel, L., *Plating and Surface Finishing*, Vol 64, No. 7, 1977, pp. 32–35.

¹¹ Proprietary chemical available from Atotech USA, Rock Hill, SC 29731.

¹² Polyvinyl chloride type lining, or integral polyvinyl type drop-in liners are available from several sources and are generally suitable for this purpose. It is advisable however to provide the supplier of the lining with the exact composition of the solution and conditions of use so proper choice of plastic and adhesive, or both can be made. Also available are preformed and or welded tanks of polyethylene and polypropylene in both normal and high density forms. These are suitable for many applications particularly if properly reinforced with external supports. Polyester fiberglass tanks may also be suitable for some applications.

Wetting agent
Current density
Time
Anode
Agitation

Optimum amount for surfactant used
250 A/m²
2 min
Nickel or inert
Work or solution movement

6.5.3 After receiving the above nickel glycolate strike, the aluminum parts can be electroplated with other metals, using standard electroplating solutions.

6.6 Zinc Immersion/Electroless Nickel Strike:

6.6.1 Aluminum parts with cleaned and conditioned surfaces are given a zinc immersion treatment as described in 6.3. A single or double immersion treatment may be used.

6.6.2 After water rinsing, the zincated parts are given an electroless nickel strike in the electroless nickel solution of choice. Because of the need for carefully buffered conditions and tolerance for dissolved zinc, these solutions are generally proprietary. The operating conditions recommended by the manufacturer should be followed carefully. In particular, it should be noted that these solutions may have deposition rates that vary with different sources, operating conditions and age. Immersion time must be adequate to ensure complete, pore-free coverage of all surfaces.

6.6.3 After receiving the electroless nickel strike, the aluminum parts can be electroplated with other metals using standard electroplating conditions, or transferred to a different electroless nickel bath for the application of electroless nickel deposits for engineering purposes.

6.7 Tin Immersion/Bronze Strike:

6.7.1 The aluminum parts should be cleaned as described in 5.1. They should then be conditioned preferably in an alkaline etch followed by rinsing and desmutting in a nitric acid plus ammonium bifluoride solution, as described in 5.2.1.

6.7.2 After water rinsing, the cleaned and conditioned aluminum parts are subjected to a tin activation treatment. This is accomplished either by simple immersion, or by “live” (current on) entry (cathodic) of the work into a proprietary aqueous stannate bath,¹¹ for 30 s at 26 to 30°C.

6.7.3 Without rinsing and with minimum time delay, the tin-activated aluminum parts are transferred into a proprietary, aqueous bronze cyanide bath¹⁰ where they are given a strike of 3 to 4 min at 26 to 30°C with a cathodic current density of 320 to 540 A/m².

6.7.4 After the bronze strike and water rinsing, other metals can be electroplated on the aluminum parts using standard electroplating solutions.

7. Safety Precautions

7.1 Some chemical solutions are exothermic upon mixing or in use, thereby requiring cooling and proper containment to prevent injury to personnel.⁴ (**Warning**—Care in the handling and use of all cyanide-containing salts and solutions must be exercised. Adequate rinsing between cyanide and acid process solutions must be performed.)



APPENDIX

(Nonmandatory Information)

X1. SOLUTIONS FOR CLEANING AND CONDITIONING ALUMINUM ALLOYS

X1.1 Carbonate-Phosphate Cleaner:

Sodium carbonate, anhydrous	25 g/L
Trisodium phosphate, anhydrous	25 g/L
Temperature	60 to 80°C
Time	1 to 3 min
Container	steel

X1.2 Deoxidizing Etch:

Sulfuric acid (density 1.83 g/mL)	100 mL
Chromic acid, CrO ₃	35 g
Water	to 1 L
Time	2 to 5 min
Temperature	70 to 80°C
Container	lined with lead

NOTE X1.1—**Warning**—Dissolve the chromic acid in approximately 800 mL of water, then slowly add the sulfuric acid with rapid mixing; when the solution has cooled to room temperature, dilute to 1 L. Fumes are toxic. Use exhaust.

X1.2.1 This solution may be used at room temperature for periods of 5 to 30 min to remove many types of oxides. Operation at this lower temperature offers greater safety and reduces the amount of hazardous fumes evolved.

X1.3 Caustic Dip:

Sodium hydroxide	50 g/L
Time	30 s to 1 min
Temperature	50°C
Container	Steel

NOTE X1.2—**Warning**—Fumes are toxic. Use exhaust.

X1.4 Acid Desmutter:

Nitric acid (density 1.4)	500 to 700 mL/L
Ammonium bifluoride	30 to 120 g/L
Time	30 s
Temperature	20 to 25°C
Container	steel with a suitable plastic lining ¹³

NOTE X1.3—**Warning**—Fumes are toxic. Use exhaust.

X1.4.1 The activity and aggressiveness of this desmutter may be controlled by varying the concentrations as indicated. Increasing the nitric acid concentration decreases activity; increasing the ammonium bifluoride concentration increases activity.

X1.5 Alternative Acid Desmutter:

Sulfuric acid (H ₂ SO ₄ 93 mass %, density 1.83 g/mL)	100 mL
Hydrogen peroxide (H ₂ O ₂ 32.5 mass %, stabilized for use with nonferrous metals)	50 mL
Water	to 1 L
Time	15 s to 1 min
Temperature	room
Container	300 series stainless steel or container with a suitable plastic lining ¹³

NOTE X1.4—**Warning**—The acid should be slowly added to 90 vol % of the water required with rapid stirring. When the solution cools to room

temperature, add the hydrogen peroxide and dilute to exact volume. Fumes are toxic. Use exhaust.

X1.6 Nitric Acid Dip:

Commercial nitric acid (67 mass %, density 1.4)	500 mL
Water	to 1 L
Temperature	room
Container	steel-lined with suitable plastic or UNS S30403, UNS S31603, or UNS S34700 stainless steel ¹³

NOTE X1.5—**Warning**—Fumes are toxic. Use exhaust.

X1.7 Sulfuric Acid Dip:

Sulfuric acid (H ₂ SO ₄ 93 mass %, density 1.83 g/mL)	150 mL
Water	to 1 L
Temperature	80°C
Time	2 to 5 min
Container	lined with lead or a suitable plastic ¹³

NOTE X1.6—**Warning**—The acid should be slowly added to the approximate amount of water required with rapid mixing. When the solution cools to room temperature, dilute to exact volume.

X1.8 Mixed Acid Dip:

Commercial nitric acid, (67 mass %, density 1.4 g/mL)	750 mL
Commercial hydrofluoric acid (48 mass %, density 1.16 g/mL)	250 mL
Time	3 to 5 s
Container	steel-lined with a suitable plastic or carbon brick or both ¹³

NOTE X1.7—**Warning**—Fumes are toxic. Use exhaust.

X1.9 Universal Deoxidizer:

Commercial nitric acid (67 mass %, density 1.4 g/mL)	500 mL
Sulfuric acid (density 1.84 g/mL)	250 mL
Water	250 mL
Ammonium bifluoride	60 g/L

NOTE X1.8—**Warning**—Add the nitric acid to the water slowly with vigorous agitation. Allow to cool to room temperature. Slowly add the sulfuric acid to the mixed acids with vigorous agitation. Allow to cool to room temperature. Dissolve required amount of ammonium bifluoride. Adjust to final volume with water if necessary. The operation may have to be interrupted several times to permit cooling! The temperature during mixing must never be allowed to exceed the safe operating limits of the lining or plastic container or irreparable damage may occur.

X1.9.1 This acid dip is applicable to almost all alloys. It is particularly useful on alloys containing magnesium.

NOTE X1.9—**Warning**—Fumes are toxic. Use exhaust.

ASTM International takes no position respecting the validity of any patent rights asserted in connection with any item mentioned in this standard. Users of this standard are expressly advised that determination of the validity of any such patent rights, and the risk of infringement of such rights, are entirely their own responsibility.

This standard is subject to revision at any time by the responsible technical committee and must be reviewed every five years and if not revised, either reapproved or withdrawn. Your comments are invited either for revision of this standard or for additional standards and should be addressed to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee, which you may attend. If you feel that your comments have not received a fair hearing you should make your views known to the ASTM Committee on Standards, at the address shown below.

This standard is copyrighted by ASTM International, 100 Barr Harbor Drive, PO Box C700, West Conshohocken, PA 19428-2959, United States. Individual reprints (single or multiple copies) of this standard may be obtained by contacting ASTM at the above address or at 610-832-9585 (phone), 610-832-9555 (fax), or service@astm.org (e-mail); or through the ASTM website (www.astm.org).